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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO	
10/089,146	09/16/2002	Wilhelm Amberg	51748	9829	
32116 759	90 05/03/2006		EXAMINER		
WOOD, PHILLIPS, KATZ, CLARK & MORTIMER			HADDAD, MAHER M		
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SUITE 3800			AKTONII	PAPER NUMBER	
CHICAGO, IL 60661			1644		
			DATE MAILED: 05/03/2004	DATE MAILED: 05/03/2006	

Please find below and/or attached an Office communication concerning this application or proceeding.

Advisory Action

Application No.	Applicant(s)	
10/089,146	AMBERG ET AL.	
Examiner	Art Unit	
Maher M. Haddad	1644	

	, , , , , , , ,					
Before the Filing of an Appeal Brief	Examiner	Art Unit				
	Maher M. Haddad	1644				
The MAILING DATE of this communication appe	ears on the cover sheet with the c	correspondence add	ress			
THE REPLY FILED <u>14 April 2006</u> FAILS TO PLACE THIS APP	LICATION IN CONDITION FOR A	LLOWANCE.				
The reply was filed after a final rejection, but prior to or on the same day as filing a Notice of Appeal. To avoid abandonment of this application, applicant must timely file one of the following replies: (1) an amendment, affidavit, or other evidence, which places the application in condition for allowance; (2) a Notice of Appeal (with appeal fee) in compliance with 37 CFR 41.31; or (3) a Request for Continued Examination (RCE) in compliance with 37 CFR 1.114. The reply must be filed within one of the following time periods:						
) Mean the period for reply expires <u>3 months from the mailing date of the final rejection.</u> The period for reply expires on: (1) the mailing date of this Advisory Action, or (2) the date set forth in the final rejection, whichever is later. In						
b) The period for reply expires on: (1) the mailing date of this A no event, however, will the statutory period for reply expire Examiner Note: If box 1 is checked, check either box (a) or TWO MONTHS OF THE FINAL REJECTION. See MPEP 7	later than SIX MONTHS from the mailin (b). ONLY CHECK BOX (b) WHEN THI	g date of the final rejecti	on.			
Extensions of time may be obtained under 37 CFR 1.136(a). The date nave been filed is the date for purposes of determining the period of example of the set forth in (b) above, if checked. Any reply received by the Office late nay reduce any earned patent term adjustment. See 37 CFR 1.704(b) NOTICE OF APPEAL	ctension and the corresponding amount shortened statutory period for reply orig r than three months after the mailing da).	of the fee. The appropr inally set in the final Offi ite of the final rejection,	iate extension fee ce action; or (2) as even if timely filed,			
The Notice of Appeal was filed on A brief in comp filing the Notice of Appeal (37 CFR 41.37(a)), or any exter a Notice of Appeal has been filed, any reply must be filed	ension thereof (37 CFR 41.37(e)), to	o avoid dismissal of th				
AMENDMENTS						
B. The proposed amendment(s) filed after a final rejection,	•		ecause			
(a) They raise new issues that would require further co		I E below);				
 (b) They raise the issue of new matter (see NOTE below); (c) They are not deemed to place the application in better form for appeal by materially reducing or simplifying the issues for appeal; and/or 						
(d) They present additional claims without canceling a NOTE: (See 37 CFR 1.116 and 41.33(a)).		jected claims.				
1. The amendments are not in compliance with 37 CFR 1.1		ompliant Amendment	(PTOL-324).			
5. Applicant's reply has overcome the following rejection(s)		•	(· · · · · · · · · · · · · · · · · · ·			
Newly proposed or amended claim(s) would be allowable if submitted in a separate, timely filed amendment canceling the non-allowable claim(s).						
7. For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is pro The status of the claim(s) is (or will be) as follows:		ill be entered and an e	explanation of			
Claim(s) allowed: <u>None</u> .						
Claim(s) objected to: <u>None</u> . Claim(s) rejected: <u>4 and 10</u> .			·			
Claim(s) withdrawn from consideration: <u>1-3 and 7-9</u> .						
AFFIDAVIT OR OTHER EVIDENCE		•				
 The affidavit or other evidence filed after a final action, be because applicant failed to provide a showing of good an was not earlier presented. See 37 CFR 1.116(e). 	ut before or on the date of filing a N nd sufficient reasons why the affidat	otice of Appeal will <u>no</u> vit or other evidence i	ot be entered s necessary and			
The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to showing a good and sufficient reasons why it is necessar	overcome all rejections under appe	al and/or appellant fa	ils to provide a			
0. The affidavit or other evidence is entered. An explanation						
REQUEST FOR RECONSIDERATION/OTHER	A done NOT whose the combination is	n namditian fan allawa				
11. The request for reconsideration has been considered by See Continuation Sheet.			nce because:			
12. Note the attached Information Disclosure Statement(s).13. Other:			1			
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	Pa	tent Examin	ا ه			



Continuation of 11. does NOT place the application in condition for allowance because:

- 1. Claim 4 stands rejected under 35 U.S.C. 103(a) as being unpatentable over Kirchengast et al (provided in the International Report and cited on the PTO-892 as reference Y) in view of Srivatsa et al (of record).
- 2. Claim 10 stands rejected under 35 U.S.C. 103(a) as being unpatentable over Kirchengast et al in view of Srivatsa et al as applied to claim 4 above, and further in view of US Pat. No. 4,761,406.

Applicant submits that the specifically claimed combination is not taught and it leads to a surprising effect. Further Applicant argues that the present invention permits the use of each compnent at a dose less than the dose useful alone, with a reduction in side effects. (page 4, lines 23-27 and page 20, lines 24-30).

However, Applicant's reliance on surprising effect do not overcome clear and convincing evidence of obviousness. Also see Richardson-Vicks Inc. v. Upjohn Co., 44 USPQ2d 1181 (CAFC 1997). The issue is whether the properties differ to such an extent that the difference is really surprising. These effects are not surprising because co-administer ETA endothelin blocker and alphavbeta3 integrin receptor antagonist is expected to provide efficacy at lower doses than the doses useful individually, with a reduction in side effects. Combination therapy are use to minimize dose-dependent side effects of an individual drug.